



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

<p>(51) International Patent Classification<sup>6</sup> : C04B 35/66, 40/06, 40/00</p>	<p>A1</p>	<p>(11) International Publication Number: <b>WO 98/52883</b> (43) International Publication Date: 26 November 1998 (26.11.98)</p>
<p>(21) International Application Number: PCT/GB98/01303 (22) International Filing Date: 18 May 1998 (18.05.98) (30) Priority Data: 9709988.1 17 May 1997 (17.05.97) GB 9727108.4 23 December 1997 (23.12.97) GB (71) Applicant (for all designated States except US): VESUVIUS K.S.R. LIMITED [GB/GB]; Sandiron House, 449/453 Abbey Lane, Beauchief, Sheffield S7 2RA (GB). (72) Inventors; and (75) Inventors/Applicants (for US only): HEY, Alan [GB/GB]; Joan Royd Cottage, Joan Royd Lane, Penistone, Sheffield S36 9DA (GB). PICKARD, David, Michael [GB/GB]; 9 Inglewood Avenue, Sothall, Beighton, Sheffield S20 2QZ (GB). (74) Agent: LUNT, Mark, George, Francis; Dibb Lupton Alsop, Fountain Precinct, Balm Green, Sheffield S1 1RZ (GB).</p>		<p>(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report.</p>
<p>(54) Title: CASTABLE REFRACTORY COMPOSITIONS</p>		
<p>(57) Abstract</p> <p>The invention relates to a method for producing refractory compositions and more particularly to a method for producing castable refractory compositions which comprises combining appropriate quantities of bulk particulate refractory aggregates substantially without pre-mixing; separately adding to the aggregate system an appropriate quantity of a pre-blended binder composition; mixing the aggregate system and the pre-blended binder composition in a mixer; and discharging the refractory composition from the mixer.</p> <div data-bbox="698 1155 1331 1827"> </div>		

**FOR THE PURPOSES OF INFORMATION ONLY**

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

## CASTABLE REFRACTORY COMPOSITIONS

This invention relates to refractory and other compositions, and more particularly to castable refractory  
5 compositions comprising particulate refractory aggregates and binder components. Compositions for iso-pressing are also within the ambit of the present invention.

Castable refractory compositions comprising particulate  
10 refractory aggregates and binder components are used in the metallurgical, cement, petrochemical and glass making industries for the production of furnace and ladle linings, launders or runners, pouring nozzles and other heat resisting applications.

15

Castable refractory compositions are generally produced by their manufacturers at sites remote from the premises of the end-users. In a conventional process for producing a castable refractory composition, raw materials, either in  
20 bulk, intermediate bulk containers, or bags on pallets comprising particulate refractory aggregates and binder components, are charged into a mixer, which is normally of around one to two tonnes capacity. With the exception of the pallets, which may be re-used, raw materials packaging  
25 is frequently subsequently discarded, especially when small paper or plastic bags are involved. Raw materials packaging is normally disposed of by land-fill or incineration, both of which are expensive processes, the former being subject

to state imposed taxes. However, residues of particulate refractory materials in the packaging also lead to waste, and therefore a reduction in this and the volume of redundant packaging which is required to be disposed of 5 would be highly desirable.

The batch of discharged raw materials is normally subjected to a thorough mixing operation in the mixer for 5-20 minutes, depending on the composition, in order to 10 produce a homogeneous composition. The power consumption of the mixer is considerable and adds to the costs of the product. Finally, the composition is discharged from the mixer and re-packaged in 25Kg, 250Kg, 500Kg, 1000Kg or 2000Kg bags as required, and after having undergone 15 extensive quality control testing, is transported to the premises of the end-user.

On arrival at the end-user's premises, the various types of bags are broken open, and the contents discharged 20 into a second mixer. The bags may again be discarded, with the attendant disposal problems set out above, and the composition is subjected to further mixing in the second mixer, during which process, water is added to the mixed composition. The castable composition is then discharged 25 from the mixer for conveying, or pumped, to the site where the cast lining is formed in-situ.

It will be apparent from the above description that there are several difficulties inherent in the current method for the manufacture of castable refractory compositions. In addition to the disposal problems 5 represented by the two sets of discarded bags, the entire composition is required to be mixed on two separate occasions, in large mixers having a substantial power consumption.

10 Having mixed aggregates with binder, it is important that the composition remains dry until the time that it is required. Otherwise the binder can deteriorate preventing homogenous curing of the final mixtures. It is for this reason, primarily, that smaller bags of the composition 15 might frequently be employed, so that if any bag does develop a leak during storage or transportation, permitting ingress of water, only a small amount of the composition is ruined.

20 However, a completely unrelated issue is that furnaces cost a considerable sum of money to be kept idle while repairs are effected. Furnace operators plan upgrades and repairs to the furnace and related equipment meticulously, so that the time for which the furnace is inoperative may 25 be minimised. One repair is the replacement of the lining of troughs, runners, furnaces and ladle linings etc, and for the refractory supplier, who usually has the responsibility of overseeing the repair, this means little

time can be allowed for the item which is to be repaired to cool before the repair must be effected so that the newly cast refractory composition will be cured and hardened before the furnace is switched back on line. Time is  
5 therefore of the essence. Furthermore, when casting refractory compositions into still very hot components which have only recently come off line, the cure rate of the composition is inevitably very fast. While this in itself is not a problem, it means that when boundaries  
10 exist between one cast and a subsequent cast, lines of weakness at the boundary can develop resulting possibly in cracks or other defects in the refractory lining.

The answer to these problems is to cast in large  
15 volumes, very quickly. Thus furnace operators may be provided with a large two-tonne mixer and a number of mobile skips or hoppers of the same capacity so that they may have the mixed composition discharged into the skips one immediately following the other. Thus, perhaps three  
20 or four skips might be filled one after the other, so that each may be discharged in rapid succession giving, to all intents and purposes, a continuous delivery. Meanwhile, the mixer is mixing a further batch, which may be ready for emptying into the first (now empty) skip or hopper, before  
25 the third skip has been discharged. In this way, perhaps up to sixteen tonnes of refractory may be cast almost at one time.

However, charging, adding water, mixing and discharging mixers is relatively time consuming, and it is this element in the process which is the bottleneck. Eventually the speed of the cycle is reduced to its slowest component and the quasi-continuous casting reduced to batch casting. Since it is the object is to have a continuous cycle of charge, mix and discharge, so that the casting process can be continuous, the mixing process needs to be speeded-up.

10

This militates against the use of small bags of composition, simply because of the time required to empty the bags into the mixer. Thus it is an object of the invention to overcome the afore-mentioned problems, or at least to mitigate their effects.

According to the present invention, the above difficulties are obviated by a novel method in which the binder composition is blended in a separate stage before mixing with the particulate refractory materials (hereinafter termed "the aggregate system").

In a first aspect, the present invention provides a method for producing castable refractory compositions which comprises composing an aggregate system of appropriate quantities of bulk particulate refractory aggregates substantially without pre-mixing, separately adding to the aggregate system an appropriate quantity of a pre-blended

binder composition, transporting the aggregate system to an installation site, mixing at the installation site the aggregate system and the pre-blended binder composition with water in a mixer and discharging the refractory composition from the mixer.

Such a method produces numerous advantages.

In a preferred embodiment of the invention, the components of the aggregate system are purchased to guaranteed chemical and particle size specification in re-usable intermediate bulk containers. The pre-blended binder is preferably added in a weighed amount to the aggregate components, and the combined materials transported to the user site in the intermediate bulk container originally containing one of the aggregate components. This has the advantage of providing at least two re-usable intermediate bulk containers, one containing the material despatched to the user site, and one available for other use.

In this specification the term "castable refractory composition" is defined as a precursor comprising particulate refractory materials, and a binder composition, which, when combined with water in the appropriate proportions preferably in the mixer, can be discharged, transported and cast in-situ to form the desired refractory shape.



The components of the aggregate system comprise refractory aggregates in the appropriate quantities, and generally comprise first and second aggregate components, 5 for example, the first comprising alumina and the second comprising silicon carbide.

The first component, alumina powder, is preferably present in an amount of from 10 to 90% of the total 10 composition by weight, and preferably has a typical particle size range of 100 to 12,500 microns.

The second component, silicon carbide is preferably present in the composition in an amount of from 5 to 90% of 15 the total composition by weight, and preferably has a typical particle size range of 5 to 2000 microns.

In each case, the particle size distribution is controlled to within a specified range.

20

Other aggregate system components may be used in whole or partial replacement for the alumina and silicon carbide components, but there will always be at least two components. Alternatives may include, for example, 25 andalusite, calcined bauxite, kyanite, sillimanite, chamottes, sintered or fused alumina-spinel, sintered or fused magnesia, zircon and zirconia.

The required particle size distribution of the aggregate components can be achieved, for example, by blending or assembling specific fractions of particulate refractory materials in the desired proportions. This operation is preferably carried out by the aggregate component suppliers who will assemble the correct combination of particles to meet a specific particle size distribution aim, and package them in a specified type and capacity intermediate bulk container.

An important aspect of the present invention is that the component parts of the binder composition are separately blended prior to batching with the aggregate system. The components of the binder composition can comprise, for example, reactive alumina, fine silicon carbide, ultrafine silica powder, clays, high alumina cements and hydrateable alumina.

The binder composition preferably comprises from 5 to 35% of the total refractory composition by weight.

The above proportions of materials refer to dry weight, and do not include the water which is added to produce the final castable refractory composition. The amount of water added can be up to about 10% by weight, but is preferably from 5 to 5.5% by weight, based upon the total dry weight of the composition.

The invention, in preferred embodiments, enables the entire composition to be mixed only once, using a mixer situated at the premises of the end-user. Moreover, it has  
5 been found that the required time of mixing is no more than has hitherto been originally required on-site when mixing water with the premixed aggregate and binder composition. In other words, substantial energy savings can be made at the refractory suppliers site by avoiding the mixing there,  
10 the saved costs of which can be passed to the final customer.

Another important advantage of the present invention is realised in that premixed aggregate and binder has very  
15 poor flow properties, due to the finer particles filling the gaps between larger particles and binding the whole together. This is, of course, very desirable in the final cast product, but not when attempts are made to transfer the components from their packaging to the mixer by solid flow.

20

Thus an important aspect of the present invention provides a package of materials for producing formable refractory products, the materials comprising, in appropriate relative quantities, substantially unmixed  
25 layers of at least first and second compositions. Preferably, said product is castable and said first and second compositions comprise aggregate components.

Preferably, said package includes a layer of a pre-blended binder composition in an appropriate quantity with respect to said aggregate components. Preferably the binder is between said first and second composition layers.

5

The components are in amounts ready to mix with water to form said castable refractory composition. This arrangement has two particular advantages. Firstly, since there is no substantial mixing between the layers, each layer is discrete and therefore flows relatively easily. This effect is enhanced if the first and second components are themselves graded into sub-layers. This arrangement enables a fast and complete discharge of the package, leaving it relatively clean for subsequent reuse.

15 Secondly, if the binder is disposed between the two aggregate layers it is protected from the two main sources of moisture ingress, namely, the open top of the package, and the bottom of the package which is susceptible to punctures and to seepage of water lying on the ground. Of course, it does not matter if the aggregates get a little damp, so long as the binder is kept dry. Since it is the binder which is the finest powder, it is usually this that gets left behind sticking to packaging walls etc. However, the aggregate sitting above the binder has the effect of

25 scrubbing the packaging as it leaves the packaging, taking with it substantially all traces of the binder.

A further advantage of this arrangement is the security it provides against deliberate attempts by others to determine the composition of the product being supplied. Precise compositions of aggregate components, and binder compositions, not to mention their total composition in the final product, are sometimes closely held trade secrets. By providing the components in segregated layers in the delivery package, an attempt to discern this information cannot successfully be made by taking only a small sample as a means of establishing the contents of the entire package.

Additionally, the present invention finds application with iso-pressable refractory compositions such as alumina/graphite, which may be formed by isostatic pressing into alumina-graphite pieces: that is to say, wherein said first and second compositions comprise alumina and graphite. The graphite may be disposed in a layer between layers of alumina.

20

In one preferred embodiment of a castable product, the components of the aggregate system, each having the required particle size distribution, are delivered to the manufacturer in re-usable packaging, which preferably comprises, for example, a mini-bulk sack.

Preferably, the alumina aggregate is delivered in an over-size mini-bulk sack which is of such a size that there

is sufficient volume for the silicon carbide and binder composition to be added to the sack. This avoids the use of paper sacks for the components, alleviating the disposal problem, and reducing the loss of components on de-bagging.

5

Of the composition components discharged into the mini-bulk sack, only the binder composition will have been subjected to prior blending. The components will, however, have been accurately weighed into the mini-bulk sack, so  
10 that they will be present in the specified proportions. The mini-bulk sack, or other suitable container, can then be delivered to the end user.

At the premises of the end-user, the mini-bulk sack is  
15 discharged directly into a mixer, and the entire refractory composition mixed for the first and only occasion. Because of the unmixed nature of the components, discharge from the sack is rapid. Water is then added to the mixer and the castable refractory composition discharged and processed in  
20 the usual way.

Power consumption is reduced, as mentioned above, since the binder represents only from 5 to 35% by weight, typically 20%, of the final mix, and this is the only  
25 component which is subjected to pre-blending.

A further advantage is that quality control testing of the components can be substantially reduced in the new

method, since every tonne of binder will typically produce 5 tonnes of final castable composition. Indeed, this latter point is particularly important because, hitherto, samples of mixed aggregate and binder could only be taken, realistically, one test for each three to six tonnes produced, depending on batch size, (representing 15-33% test rate) so that quality could not be so well assured. With the present method, however, the aggregates are supplied by the aggregate supplier in guaranteed composition having been quality assured separately. The refractory supplier need only be concerned with the binder for which a single batch, comprising typically 10 to 20% by weight of the final composition, will typically be sufficient for, say, six tonnes of final product. Testing of the premixed binder therefore represents 100% quality assurance.

The method according to the invention also enables re-usable packaging to be introduced, thereby reducing the problem of disposal of unwanted packaging materials.

Overall, there is less raw material handling and therefore reduced waste/spillage of raw materials. The number of weighing operations can also be reduced and overall batch accuracy can thereby be improved.

25

Whilst it is usual for the manufacturer to supply the refractory composition to the end-user in bulk re-usable containers ready for mixing, it would be possible for the

aggregate system and the blended binder composition to be delivered to the end-user separately and discharged into a mixer at the end-user's premises.

5        Additionally, the present invention is concerned with iso-pressing compositions such as alumina/graphite, which may be isopressed into alumina-graphite pieces. The advantages of the present invention may likewise be felt with such compositions.

10

A comparison between the conventional processing method and a method in accordance with the invention, is illustrated, by way of example, in the accompanying flow-diagrams, in which:

15

Diagram 1 is a schematic representation of a conventional method for producing castable refractory compositions; and

Diagram 2 is a schematic representation of a method  
20 for producing castable refractory compositions according to the invention.

In Diagrams 1 and 2, the essential difference between the two processes is the mixing station 12 in Diagram 1 is  
25 applied to all materials, whereas the same station 12' in Diagram 2 only relates to blending of the binder. Moreover, there is an additional input 14 in Diagram 2 in which pre-weighed aggregates, certified by their supplier



to conform with predetermined specifications, are supplied direct to the batching station 16.

The invention is further described hereinafter, by way of example, with reference to the accompanying drawing, in which is a section through a mini-bulk sack showing the layering of the materials contained therein.

In the drawing, a minibulk sack 16 comprises a reinforced fabric bag having carrying handles 18 and discharge opening 20. The opening 20 is controlled by a tie 22. In the bag there are three layers of material: a first layer 24 of a first aggregate component such as alumina; a second layer 26 of a binder composition; and, a third layer 28 of a second aggregate component such as silicon carbide. The binder layer 26 is a pre-blended, homogenous composition, such as described below. However, the aggregate layers 24, 28 may be stratified according to its different components with substantially no inter-mixing.

In any event, there is little or no intermixing between the three layers, other than marginal mixing and mutual settlement at their respective boundaries. Importantly there is no opportunity for mutual interpacking of the various components, so that smaller particles of the components do not fill and pack the interstices between the larger particles.

An embodiment of the invention will now be illustrated by the following Examples:

**EXAMPLE I**

This example describes the production of a castable refractory composition by a method in accordance with the invention.

10      Binder Preparation

The component parts of a binder composition comprising ultrafine alumina powder, silicon carbide powder, high alumina cements, clays and deflocculants are charged into a 15 suitable mixer, capable of producing a large batch.

Packaging from the binder ingredients is discarded. On completion of the mixing cycle, the binder composition is discharged from the mixer into suitable containers or packaging which may include re-usable items. At this stage, a sample of the binder composition is taken and blended with the correct proportion of the aggregate system to give a complete mix for quality control testing in the laboratory. The physical properties of the final composition are then determined and compared to the product definition or specification.

### Aggregate System Preparation

1.0 tonne of alumina aggregate of average particle size range 12500 to 100 microns is provided in a mini-bulk sack of total capacity 2.0 tonnes. 0.6 tonnes of silicon carbide aggregate of particle size range 2000 to 5 microns is provided in a mini-bulk sack of capacity 1.0 tonne.

Additional aggregate materials, for example, sintered or fused alumina spinel may be added to bring the aggregate system to the desired specification. Alternatively, the sintered or fused alumina-spinel aggregates may wholly or partially replace the alumina component. In any event, the components of each aggregate are added separately so that, in each sack, the aggregates are in sub-layers within the sack.

0.4 tonnes of mixed binder composition are added to the mini-bulk sack containing the alumina aggregate component, followed by the 0.6 tonnes of the silicon carbide aggregate component, to produce a package of layered components. Of course, pouring the sub-layered silicon carbide component into the alumina sack does result in some mixing, but not enough to homogenise that component.

The sack is then sealed for transportation to the end-user site of installation.

At the premises of the end-user, the entire contents of the mini-bulk sack are discharged into a mixer of capacity 2 tonnes. By virtue of the layering of the components, discharge takes about 15 seconds, as opposed to about 30 5 seconds with mixed components. Moreover, the discharge is clean, leaving the sack reusable. The components are mixed for 2 minutes in the mixer while typically 100 litres of water are added, and, after further mixing for about 4 minutes to distribute the water evenly throughout the 10 composition, the castable refractory composition is discharged into appropriate conveying systems for transporting to the area where the castable composition is to be installed.

15

**EXAMPLES A to D**

Three further examples (A, B and C) were prepared for comparison with a fourth, conventionally premixed, example (D). All Examples A to D have identical compositions, and each Example A to C was dry mixed for reducing periods of 20 time before being mixed with water.

The following Table of results was achieved:-

TABLE

	Examples			
	A	B	C	D
<b>Mixing cycle used</b>				
Dry mixing (min)	7.0	4.0	0.5	0.5
Add water (min)	1.5	1.5	1.5	1.5
Wet mixing (min)	4.0	4.0	4.0	4.0
TOTAL TIME (min)	12.5	9.5	6.0	6.0
<b>Physical Properties</b>				
Flow value (mm)	154	159	157	150
Setting time (min)	160	185	190	180
<b>Dried Properties</b> (110°C)				
Bulk Density (kg m <sup>3</sup> )	2820	2830	2840	2850
CCS (MN m <sup>2</sup> )	23.6	28.5	27.6	25.2

Example A represents an equivalent mixing time to the conventional Example D, given that example D will have been subjected to pre-mixing for about six minutes at the refractory supplier's premises before delivery to the installation site. Example C, however, shows that reducing the dry mixing stage to no more than was conventionally done, prior to addition of water, has no significant effect on the physical and dried properties of the final product compared with Example D, despite reduced mixing effected.

CLAIMS

1. A method for producing castable refractory compositions, which method comprises:-
  - 5 composing an aggregate system of appropriate quantities of bulk particulate refractory aggregates substantially without pre-mixing;  
separately adding to the aggregate system an appropriate quantity of a pre-blended binder composition;  
10 transporting the aggregate system to an installation site;  
mixing at the installation site the aggregate system and the pre-blended binder composition with water in a mixer; and,  
15 discharging the refractory composition from the mixer.
2. A method according to Claim 1, wherein the aggregate system and binder composition are transported in  
20 a re-usable bulk container.
3. A method according to Claim 2, in which the re-usable bulk container is a mini-bulk sack.
- 25 4. A method according to Claim 1, in which the aggregate system and the pre-blended binder composition are discharged directly into the mixer and mixed therein.

5. A method according to any of the preceding claims, in which the aggregate system comprises first and second components.

5

6. A method as claimed in claim 5, in which said first and second components are alumina and silicon carbide respectively.

10 7. A method according to Claim 5 or 6, when dependent on claim 3, in which the first aggregate component is delivered in said over-size mini-bulk sack which is of such a size that there is sufficient volume for the second aggregate component and binder composition to be  
15 added to the sack.

8. A method according to any of the preceding claims, in which the entire refractory composition is mixed only once.

20

9. A method according to any of the preceding claims, substantially as described in the Examples.

10. A method according to any of the preceding claims  
25 substantially as hereinbefore described.

11. A particulate refractory composition produced by a method according to any of the preceding claims.

12. A castable refractory composition produced by a method according to any of the preceding claims.

13. A package of materials for producing formable refractory products, the materials comprising, in appropriate relative quantities, substantially unmixed layers of at least first and second compositions.

14. A package as claimed in claim 13, in which said product is castable and said first and second compositions comprise aggregate components.

15. A package as claimed in Claim 14, in which said package includes a layer of a pre-blended binder composition, in an appropriate quantity with respect to said aggregate components.

16. A package as claimed in Claim 15, in which the binder is in a layer between said first and second composition layers, whereby said binder is at least partially protected from ingress of moisture into the package from either side of the first and second composition layers remote from said binder layer.

17. A package as claimed in any of Claims 14 to 16, in which said first composition is alumina, and said second component is silicon carbide.



18. A package as claimed in any of Claims 14 to 16,  
in which said first and second components are the same.

5 19. A package as claimed in claim 13, in which said  
product is iso-pressable, and said first and second  
compositions comprise alumina and graphite.

20. A package, substantially as hereinbefore  
10 described with reference to the Examples.

1/3

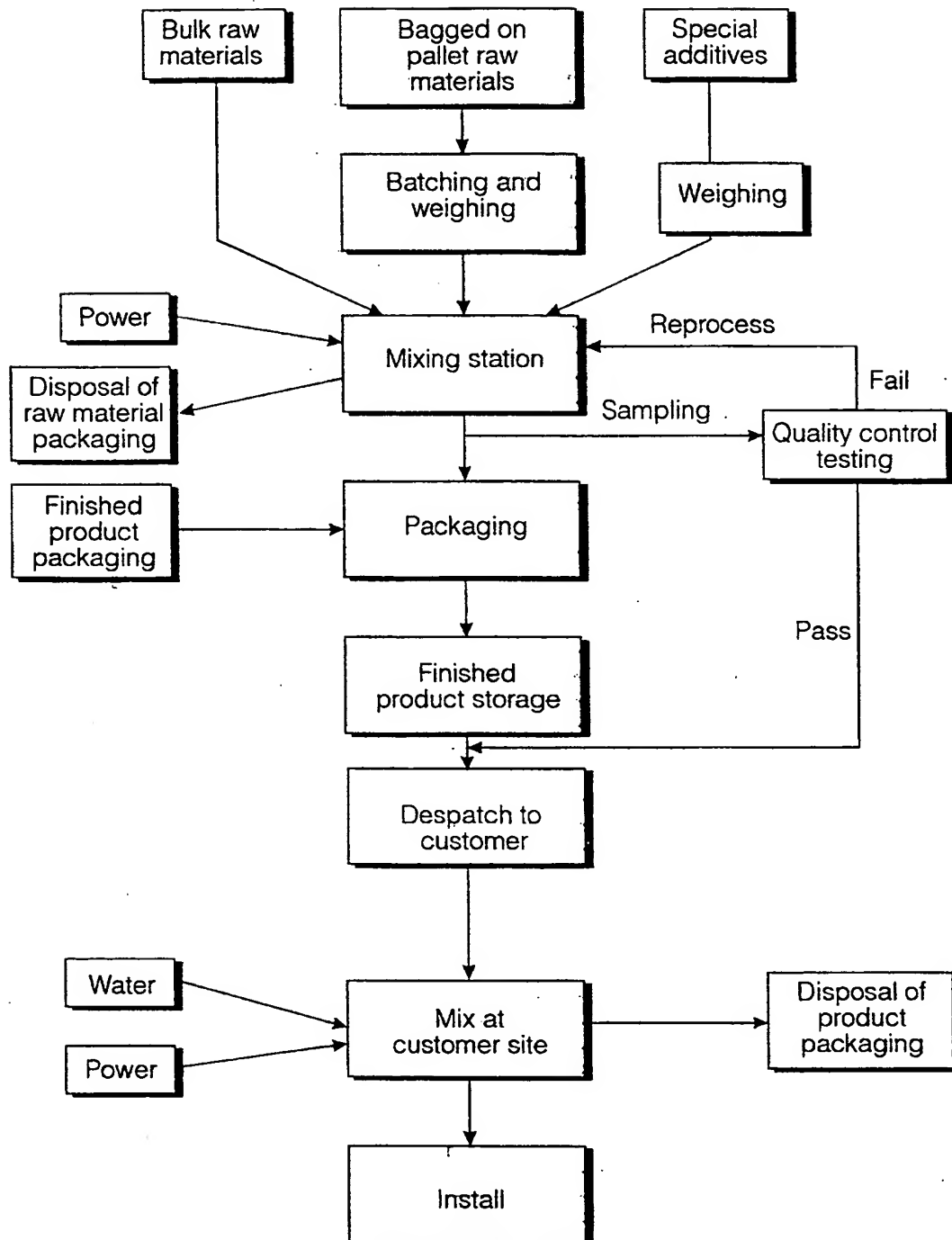


Diagram 1

2/3

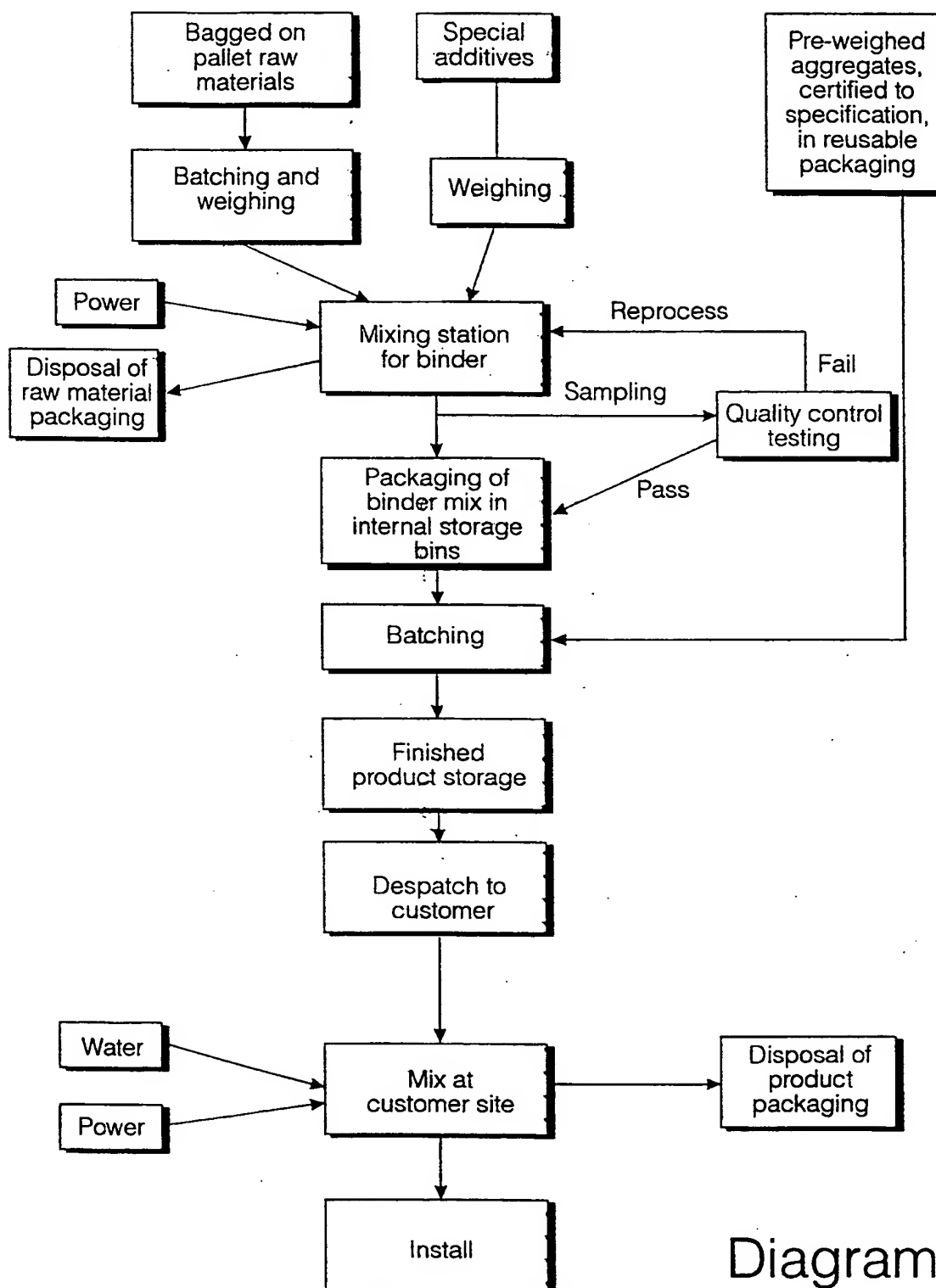


Diagram 2

3/3

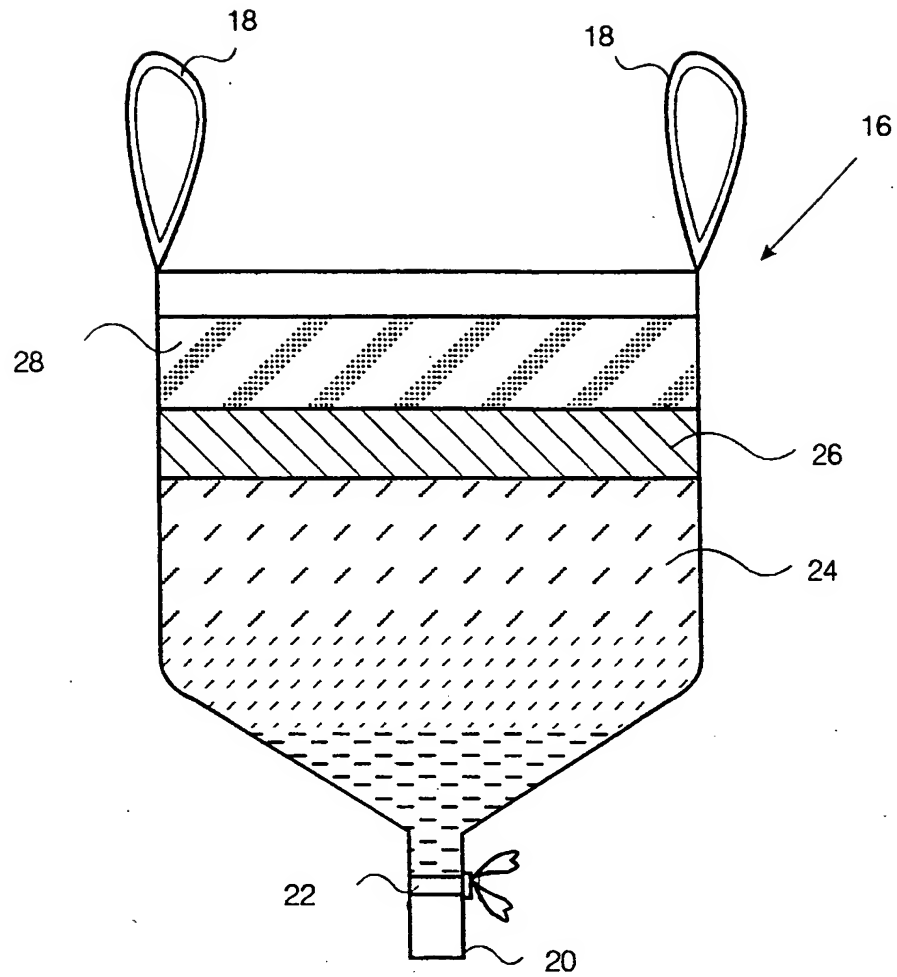


Fig. 1

# INTERNATIONAL SEARCH REPORT

International Application No

PCT/GB 98/01303

## A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C04B35/66 C04B40/06 C04B40/00

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C04B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 494 267 A (ANDERSON MICHAEL W ET AL) 27 February 1996 see examples 1,2	11,12
A	---	1
X	DATABASE WPI Section Ch, Week 8712 Derwent Publications Ltd., London, GB: Class L02, AN 87-084047 XP002071414 & JP 62 036 071 A (SHINAGAWA FIRE BRICK) see abstract	11,12
A	---	1
	--- -/--	

☒ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

### Special categories of cited documents:

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "3" document member of the same patent family

Date of the actual completion of the international search

14 July 1998

Date of mailing of the international search report

23/07/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel: (+31-70) 340-2040. Tx. 31 651 epo nl.  
Fax: (+31-70) 340-3016

Authorized officer

Mini, A

# INTERNATIONAL SEARCH REPORT

national Application No  
PCT/GB 98/01303

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0 579 994 A (TAIKOROZAI CO LTD) 26 January 1994 see examples 1-5 see page 3, line 25 - line 30	11,12
A	see page 2, line 26 - line 29 ----	1
A	EP 0 752 397 A (ASAHI GLASS CO LTD) 8 January 1997 see claims 1,7,8 ----	1
A	US 4 174 226 A (ALEXANDERSON WALTER H ET AL) 13 November 1979 -----	

# INTERNATIONAL SEARCH REPORT

Information on patent family members

national Application No

PCT/GB 98/01303

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5494267 A	27-02-1996	NONE	
EP 0579994 A	26-01-1994	JP 2556418 B	20-11-1996
		JP 6048845 A	22-02-1994
		DE 69300760 D	14-12-1995
		DE 69300760 T	15-05-1996
		US 5589426 A	31-12-1996
EP 0752397 A	08-01-1997	JP 9071478 A	18-03-1997
US 4174226 A	13-11-1979	AR 216340 A	14-12-1979
		AU 501778 A	28-06-1979
		BR 7805130 A	10-04-1979
		CA 1093103 A	06-01-1981
		DE 2839080 A	22-03-1979
		DK 387178 A	16-03-1979
		FR 2403315 A	13-04-1979
		GB 2004266 A, B	28-03-1979
		JP 1189445 C	13-02-1984
		JP 54053121 A	26-04-1979
		JP 58020908 B	26-04-1983
		SE 7808685 A	16-03-1979